Table 18: Clinical evidence profile: vitamin C versus allopurinol (in people not taking allopurinol at baseline)

Nº of studies Study design Risk of bias Inconsistency Indirectness Imprecision Considerations Considerations Considerations Studies S	Certainty assessment № of patients									atients	Effe	ct		
					Inconsistency	Indirectness	Imprecision			(people not taking allopurinol	Relative		Certainty	Importance

1	randomised	very	not serious	not serious	not serious	none	10	10	-	MD <b>0.15</b>	ФФОО	CRITICAL
	trials	serious <sup>a</sup>								higher	LOW	
										(0.08		
										higher to		
										0.22		
										higher)		
										,		

a. Downgraded by 1 increment if the majority of the evidence was at high risk of bias, and downgraded by 2 increments if the majority of the evidence was at very high risk of bias. For dichotomous outcomes MIDs were taken to be RRs of 0.8 and 1.25. For continuous outcomes 0.5 x baseline SD for continuous outcomes. Where there was only one study 0.5 (SMD) was used for continuous outcomes.